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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/769,404	01/26/2001	Theo Wallmann	8932-296	4809
20582	7590	09/20/2005	EXAMINER	
JONES DAY 51 Louisiana Avenue, N.W. WASHINGTON, DC 20001-2113			WANG, SHENGJUN	
		ART UNIT		PAPER NUMBER
		1617		

DATE MAILED: 09/20/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

<b>Office Action Summary</b>	Application No.	Applicant(s)
	09/769,404	WALLIMANN ET AL.
	Examiner	Art Unit
	Shengjun Wang	1617

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

**Period for Reply**

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

**Status**

- 1) Responsive to communication(s) filed on 13 June 2005.
- 2a) This action is FINAL.                            2b) This action is non-final.
- 3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

**Disposition of Claims**

- 4) Claim(s) 1-6,8-18 and 21-27 is/are pending in the application.
- 4a) Of the above claim(s) 4-6,8-12,15-18 and 25-27 is/are withdrawn from consideration.
- 5) Claim(s) \_\_\_\_\_ is/are allowed.
- 6) Claim(s) 1-3,13,14,21-24 is/are rejected.
- 7) Claim(s) \_\_\_\_\_ is/are objected to.
- 8) Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

**Application Papers**

- 9) The specification is objected to by the Examiner.
- 10) The drawing(s) filed on \_\_\_\_\_ is/are: a) accepted or b) objected to by the Examiner.  
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).  
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

**Priority under 35 U.S.C. § 119**

- 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) All    b) Some \* c) None of:
  1. Certified copies of the priority documents have been received.
  2. Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
  3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

**Attachment(s)**

- 1) Notice of References Cited (PTO-892)
- 2) Notice of Draftsperson's Patent Drawing Review (PTO-948)
- 3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)  
Paper No(s)/Mail Date \_\_\_\_\_
- 4) Interview Summary (PTO-413)  
Paper No(s)/Mail Date. \_\_\_\_\_
- 5) Notice of Informal Patent Application (PTO-152)
- 6) Other: \_\_\_\_\_

*kc*

*Scd*

## DETAILED ACTION

Receipt of applicants' amendments and remarks submitted June 13, 2005 is acknowledged.

### *Claim Rejections 35 U.S.C. 103*

1. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

2. Claims 1-3, 13, 14, 21-24 are rejected under 35 U.S.C. 103(a) as being unpatentable over Kaddurah-Daouk (US 5,998,457) in view of Meisner (US 4,772,591), Grant et al. (US 5,888,553), Beale (US 5,756,469) and Beale (US 5,716,926).

3. Kaddurah-Daouk teaches a method of treating osteoporosis or osteoarthritis comprising administering therapeutical effective amount of creatine compound, or a pharmaceutical acceptable salt, to patient. See, particularly, the abstract, table 1-2, and claims 1-12.

4. Kaddurah-Daouk does not teach expressly the employment of creatine pyruvate for the treatment, or the particular amount administered, or the method may be employed for promoting growth and mineralization of bone; improving acceptance and osseous integration of bone; or accelerating healing as claimed in claims 22-24, or the purity as herein required.

5. However, Grant et al. teaches that the excess of cortisol is known to be a cause of osteoporosis, tissue degeneration, and an anabolic composition with anticortisol effect are used to balance effect of cortisol. The anabolic composition comprising creatine. See, column 1, line

52 bridging column 2, line 59, column 5, lines 56-65 and claim 8. Meisner teaches a method for accelerated wound healing or treating degenerative disorders including periodontal disease osteoarthritis, comprising administering a composition comprising creatine to an animal or human. See, particularly, column 1, line 28 bridging column 2, line 45, column 5, lines 3 bridging column 7, line 10. Beale ('469) teaches creatine pyruvate (pyruvyl-creatine) is particularly useful as cortisol antagonist or cortisol blocker for prevent the catabolic activity of cortisol. See column 1, lines 7-18, 54-60; column 3, lines 46-63, and column 5, lines 54-60. Beale ('926) further teaches that pyruvate is known to be useful for treating osteoporosis. See, claim 24. Furthermore, it is noted that none of the cited references require the presence of all of the three compounds excluded herein: dihydrotriazine, dicyano-diamide, and creatinine.

Therefore, it would have been *prima facie* obvious to a person of ordinary skill in the art, at the time the claimed the invention was made, to employ creatine pyruvate composition, which is essentially free at least one of the three compounds, for treating connective tissue degenerative disorders, including those unrelated to weight gain or weight lose, such as osteoporosis, osteoarthritis or periodontitis, or for accelerating wound healing, promoting growth of connective tissue (cartilage). Note claims 23 and 24 read on the composition comprising creatine pyruvate, since creatine pyruvate is both a creatine salt, and a pyruvate.

A person of ordinary skill in the art would have been motivated to employ creatine pyruvate for treating connective tissue degenerative disorders, including those unrelated to weight gain or weight lose, such as osteoporosis, osteoarthritis or periodontitis, or for accelerating wound healing, promoting growth of connective tissue (cartilage) because it is *prima facie* obvious to combine two compounds each of which is taught in the prior art to be

useful for same purpose in order to form third composition that is to be used for very the same purpose; idea of combining them flows logically from their having been individually taught in prior art; thus , the claimed invention which employ a combination (salt) of two compounds known to be useful for treating osteoporosis sets forth *prima facie* obvious subject matter. See In re Kerkhoven, 205 USPQ 1069. Note treating osteoporosis is to promoting mineralization of bone. Further, creatine pyruvate is particularly known to be useful for treating disease associated with cortisol activity and connective tissue degenerative disorders is known to be closely related to cortisol activity. Claims 22-24 are obvious because creatine is known to be useful for promoting tissue repair process, and treating osteoarthritis and osteoporosis would also considered as a process of promoting tissue (cartilage) repairing since one of the major symptoms of osteoarthritis and osteoporosis is tissue degeneration. As to claims 28 and 30, note, in view of the teachings of Beale, Meisner and Grant, one of ordinary skill in the art would have appreciated that therapeutic effect of creatine pyruvate is not limited only to the symptoms related to weight gain or weight lose. Claims 13 and 14 are interpreted broadly as read on the elected invention, i.e., no foreign tissue have been introduced into the bond, since human bone are known to contain cells in general and chondroblasts cell particular. Finally, The optimization of a result effective parameter, e.g., the effective amount of creatine, is considered within the skill of the artisan. See, In re Boesch and Slaney (CCPA) 204 USPQ 215. Furthermore, the instant claims are essentially directed to a particular salt of creatine for treating disorders known to be treated by creatine, or its derivatives, or its salts. Absent evidence to the contrary, the employment of pyruvate creatine is seen to be a selection from amongst equally suitable material and as such obvious. Ex parte Winters 11 USPQ 2<sup>nd</sup> 1387 (at 1388). As to the negative limitation,

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"essentially free of one or more of dihydrotriazine, dicyano-diamide, or creatinine," note since the cited references do not teach or suggest the particular requirement of the three compounds, the suggested method would encompass the employment of a composition essentially free of at least one of the compounds. What is claimed herein is a specific range (essentially free) within a broad range (not required). It is well settled that in the case where the claimed ranges "overlap or lie inside ranges disclosed by the prior art" a *prima facie* case of obviousness exists. *In re Wertheim*, 541 F.2d 257, 191USPQ 90 (CCPA 1976); *In re Woodruff*, 919 F.2d 1575, 16 USPQ2d 1934 (Fed.Cir. 1990).

*Response to the Arguments*

Applicants' amendments and remarks submitted June 13, 2005 have been fully considered, but are unpersuasive.

Applicants contend that the cited references do not teach or suggest the particular daily amount of creatine pyruvate herein since the cited references disclose much large daily dosage of creatine (Kaddurah-Daouket al.) or pyruvate creatine (Beale et al). The arguments are found unpersuasive.

6. As discussed in the prior office action, one cannot show nonobviousness by attacking references individually where the rejections are based on combinations of references. See *In re Keller*, 642 F.2d 413, 208 USPQ 871 (CCPA 1981); *In re Merck & Co.*, 800 F.2d 1091, 231 USPQ 375 (Fed. Cir. 1986). Particularly, considering the cited references as a whole, it is known that: creatine compounds are useful for treating osteoporosis or osteoarthritis; excess of cortisol is a cause of osteoporosis, tissue degeneration, and anabolic composition with anticortisol effect are useful for balancing the effect of cortisol; creatine pyruvate is useful for accelerating would

healing or for treating degenerative disorders, including osteoarthritis; and creatine pyruvate (the elected compound, a salt of creatine) is a cortisol antagonist. Possessing all the knowledge mentioned above, one of ordinary skill in the art, would have seen the claimed invention, which is drawn to the employment of creatine pyruvate for treating bond or cartilage condition, or for promoting healing of bone disorders, *prima facie* obvious. The effective amount of creatine for treatment of the particular bond disorders would be a matter of optimization of results affecting parameters, which is within the skill of artisan. Further, Kaddurah-Daouket do not limit the amount of creatine to the large amount as recited by applicants. To the contrary, Kaddurah-Daouket states “For this invention (including treating osteoporosis and osteoarthritis) the creatine compound will be administered at dosage and for periods of time effective to reduce, ameliorate or eliminate the symptoms of the disease. Dose regimens may be adjusted for purpose of improving the therapeutic or prophylactic response of the compound.” (column 11, lines 36-43). Therefore, optimization of the effective amounts for treatment of the particular conditions would have been obvious to one of ordinary skill in the art. It is further noted the amounts recited by applicants in the response are not particularly designed for treatment of the disorders herein listed.

7. **THIS ACTION IS MADE FINAL.** Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period

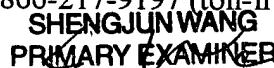
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will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Shengjun Wang whose telephone number is (571) 272-0632. The examiner can normally be reached on Monday to Friday from 7:00 am to 3:30 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Sreeni Padmanabhan, can be reached on (571) 272-0629. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

  
SHENGJUN WANG  
PRIMARY EXAMINER

Shengjun Wang  
Primary Examiner  
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